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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/560,933	06/14/2006	Giuseppe Zattera	82062-0187	4073
24633	7590	08/22/2008	EXAMINER	
HOGAN & HARTSON LLP IP GROUP, COLUMBIA SQUARE 555 THIRTEENTH STREET, N.W. WASHINGTON, DC 20004			PATEL, SHEFALI DILIP	
			ART UNIT	PAPER NUMBER
			3767	
			NOTIFICATION DATE	DELIVERY MODE
			08/22/2008	ELECTRONIC

**Please find below and/or attached an Office communication concerning this application or proceeding.**

The time period for reply, if any, is set in the attached communication.

Notice of the Office communication was sent electronically on above-indicated "Notification Date" to the following e-mail address(es):

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<b>Office Action Summary</b>	<b>Application No.</b>	<b>Applicant(s)</b>	
	10/560,933	ZATTERA, GIUSEPPE	
	<b>Examiner</b>	<b>Art Unit</b>	
	SHEFALI D. PATEL	3767	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

#### Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

#### Status

1) Responsive to communication(s) filed on 12/15/2005, 06/14/2006, and 07/30/2008.

2a) This action is **FINAL**.                                   2b) This action is non-final.

3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

#### Disposition of Claims

4) Claim(s) 1-15 and 18-24 is/are pending in the application.

4a) Of the above claim(s) 22 and 23 is/are withdrawn from consideration.

5) Claim(s) \_\_\_\_\_ is/are allowed.

6) Claim(s) 1-15, 18-21 and 24 is/are rejected.

7) Claim(s) \_\_\_\_\_ is/are objected to.

8) Claim(s) \_\_\_\_\_ are subject to restriction and/or election requirement.

#### Application Papers

9) The specification is objected to by the Examiner.

10) The drawing(s) filed on 15 December 2005 is/are: a) accepted or b) objected to by the Examiner. Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a). Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).

11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

#### Priority under 35 U.S.C. § 119

12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).

a) All    b) Some \* c) None of:

- Certified copies of the priority documents have been received.
- Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
- Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\* See the attached detailed Office action for a list of the certified copies not received.

#### Attachment(s)

1) Notice of References Cited (PTO-892)

2) Notice of Draftsperson's Patent Drawing Review (PTO-948)

3) Information Disclosure Statement(s) (PTO/SB/08)  
Paper No(s)/Mail Date 06/14/2006.

4) Interview Summary (PTO-413)  
Paper No(s)/Mail Date. \_\_\_\_\_.

5) Notice of Informal Patent Application

6) Other: \_\_\_\_\_.

## **DETAILED ACTION**

### ***Election/Restrictions***

1. Claims 22 and 23 are withdrawn from further consideration pursuant to 37 CFR 1.142(b) as being drawn to a nonelected Group II, there being no allowable generic or linking claim. Election was made **without** traverse in the reply filed on July 30, 2008.
2. Currently, claims 1-15, 18-21, and 24 are under examination, based on the election of Group I.

### ***Specification***

3. Applicant is reminded of the proper language and format for an abstract of the disclosure: The abstract should be in narrative form and generally limited to a single paragraph on a separate sheet within the range of 50 to 150 words. It is important that the abstract not exceed 150 words in length since the space provided for the abstract on the computer tape used by the printer is limited. **The form and legal phraseology often used in patent claims, such as "means" and "said," should be avoided.** The abstract should describe the disclosure sufficiently to assist readers in deciding whether there is a need for consulting the full patent text for details.

The language should be clear and concise and should not repeat information given in the title. It should avoid using phrases which can be implied, such as, "The disclosure concerns," "The disclosure defined by this invention," "The disclosure describes," etc.

In regards to the Abstract, the usage of "first and second occluding **means** suitable for" should be corrected to eliminate the usage of "**means**".

### ***Claim Objections***

4. Claims 1 and 24 are objected to because of the following informalities:

In regards to claims 1 and 24, the openings [24] are used by two terms in the claims: “at least one opening” and “openings”. To avoid confusion, one term should be consistently used within the claims.

Appropriate correction is required.

***Claim Rejections - 35 USC § 103***

5. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

6. Claims 1-7, 12-15, 18, 20, 21, and 24 are rejected under 35 U.S.C. 103(a) as being unpatentable over Frazee et al (US 5,908,407), and further in view of Don Michael (US 5,163,905).

In regards to claims 1-3 and 24, Frazee et al teaches a catheter (catheter [10]) for medical applications, suitable for being inserted into a duct comprising a first vessel (left transverse sinus [72] to right transverse sinus [74]) and a second vessel (superior sagittal sinus [56]) which branches off from said first vessel (Figure 8), the catheter [10] (Figure 4) comprising:

a. a catheter body (elongate tube [90]) which extends from a proximal end (proximal end [30]) to a distal end (distal end [32]), said catheter body [90] comprising a main cavity (through lumen [101]), bounded by a lateral wall, which passes through the catheter body [90] between the proximal end [30] and the distal end [32] (Figure 5)(column 4, lines 59-62), suitable for receiving a guide cable (guidewire [18]) for the

insertion of the catheter into the first vessel (column 5, lines 20-28)(Figure 8), and at least one opening (ports [114][116][118]), disposed on the lateral wall at the distal end [32] and suitable for perfusing a substance (Figure 4)(column 5, lines 20-28), characterized in that the catheter body [90], at a portion of the lateral wall comprised between said at least one opening [114][116][118] and said distal end [32], comprises:

i. first (occlusion balloon [41]) (Figure 4) and second occluding means (septum valve [127]) (Figure 7), wherein the first occluding means [41] are suitable for at least partially occluding a gap between the catheter body [90] and an inner wall of the first vessel [72][74] (Figure 8), and the second occluding means [127] can be associated internally with said main cavity [101] and are suitable for at least partially occluding said main cavity (column 5, lines 29-31)

ii. said first [41] and second occluding means [127] defining a preferred direction of outflow (flow [62]) of a fluid from the main cavity [101] of the catheter body [90] to the second vessel [56], through said at least one opening [114][116][118] of the catheter body (Figure 8)

b. wherein all the openings [114][116][118] pass through said lateral wall and are in fluid communication with the main cavity [101] (column 5, lines 6-10)

c. said at least one opening [114][116][118] is such that the area of the at least one opening (Figure 4) is not less than the area of the cavity (at hole [125] of through lumen [101]) of the distal end [32] of the catheter body [90] (Figure 7)

Frazee et al does not teach that said openings [114][116][118] are not aligned with one another with respect to a main axis of extension of the catheter body [90], since Frazee et al teaches that

said openings are aligned with each other with respect to the main axis of the catheter body (Figure 4). Don Michael teaches a catheter (catheter [2]) comprising openings (openings [28][30]), wherein said openings are not aligned with the main axis of the catheter, since said openings are disposed in a helical fashion about the catheter (Figure 2). It would have been obvious to a person having ordinary skill in the art at the time the invention was made to modify the catheter body of Frazee et al with non-aligned openings, as taught by Don Michael, as an obvious design choice to the user, since it has been held that rearranging parts (openings) of an invention involves only routine skill in the art. *In re Japikse*, 86 USPQ 70. Also, non-aligned openings will provide a wider area and direction of fluid flow, since the location of the openings is not restricted to one axis.

*From Applicant's specification, the first occluding means is an inflatable element (page 8, lines 19-20), and the second occluding means is an occluding body and an insertion cable (page 9, lines 14-17).*

In regards to claim 4, in a modified catheter of Frazee et al and Don Michael, Frazee et al teaches that said first occluding means [41] comprises an inflatable element positioned round the catheter body [90] (Figure 4), said inflatable element [41], in a rest state, adhering substantially to the catheter body [90], and in a working state, being substantially in contact with the inner wall of the first vessel [72][74] (Figure 8) (column 3, lines 58-63).

In regards to claim 5, in a modified catheter of Frazee et al and Don Michael, Frazee et al teaches that said inflatable element [41] is in fluid communication with the proximal end [30] so as to be operable from said proximal end (column 4, lines 62-67).

In regards to claim 6, in a modified catheter of Frazee et al and Don Michael, Frazee et al teaches that said catheter body [90] comprises a secondary cavity (inflation lumen [105]), which extends from the proximal end [30] to the distal end [32] and is hermetically separated from said main cavity [101] (Figure 5), said secondary cavity [105] being in fluid connection with said first occluding means [41] so as to permit the actuation of said first occluding means (column 4, lines 59-67).

In regards to claim 7, in a modified catheter of Frazee et al and Don Michael, Frazee et al teaches that said secondary cavity [105] is produced in a thickness of said lateral wall of said catheter body [90] (Figure 5).

In regards to claim 12, in a modified catheter of Frazee et al and Don Michael, Frazee et al teaches that said catheter body [90], at said distal end [32], comprises a portion with a tapered profile so as to reduce the cavity of the catheter body at the distal end (Figure 7).

In regards to claim 13, in a modified catheter of Frazee et al and Don Michael, Frazee et al teaches that said second occluding means, at said distal end [32], comprises a membrane [127] suitable for at least partially occluding said main cavity [101] and having a hole (slit [130]) suitable for allowing the passage of the guide cable [18] of the catheter (Figure 7) (column 5, lines 29-35).

In regards to claim 14, in a modified catheter of Frazee et al and Don Michael, Frazee et al teaches that said membrane [127] is firmly connected to the distal end [32] of the catheter body [90] (Figure 7) (column 5, lines 29-31).

In regards to claim 15, in a modified catheter of Frazee et al and Don Michael, Frazee et al does not state that the membrane [127] is made of a material suitable for being sterilized. However, it would have been obvious to a person having ordinary skill in the art at the time the invention was made to provide a sterilizable material for the membrane, since it was known in the art, as common practice in the art, to sterilize medical equipment, such as catheters, in order to eliminate transmissible agents (bacteria, viruses, etc.) from medical surfaces in order to prevent contamination to the environment.

In regards to claim 18, in a modified catheter of Frazee et al and Don Michael, Frazee et al teaches a main pathway (connector [103]), at said proximal end [30], that is suitable for receiving said second occluding means [127] and is fluidly connected to said main cavity [101] (Figure 4) (column 4, lines 59-62).

In regards to claim 20, in a modified catheter of Frazee et al and Don Michael, Frazee et al teaches that said proximal end [30] comprises a secondary pathway (connector [107]), fluidly connected to said secondary pathway [105], and suitable for receiving at the inlet a fluid, so as to allow the flow of the fluid from the proximal end [30] to the distal end [32] (Figure 4) (column 4, lines 62-67).

In regards to claim 21, , in a modified catheter of Frazee et al and Don Michael, Frazee et al teaches that said proximal end [30] comprises am infusion pathway (connector [103]), fluidly connected to said main cavity [101] and suitable for receiving at the inlet a fluid, so as to allow

the flow of the fluid from the proximal end [30] to the distal end [32] (Figure 4) (column 4, lines 59-62)(column 5, lines 20-28).

7. Claim 8 is rejected under 35 U.S.C. 103(a) as being unpatentable over Frazee et al and Don Michael, as applied to claim 6 above, and further in view of Prosl (US 5,868,717).

In regards to claim 8, in a modified catheter of Frazee et al and Don Michael, Frazee et al does not teach that said catheter body [90] has an oval cross-section with a first pole more pronounced than a second pole, since Frazee et al teaches that said catheter body has a circular cross-section (Figure 5). Prosl teaches a catheter [10] having an oval cross-section, wherein a first pole (second wall [30]) is more pronounced than a second pole (first wall [20] diametrically opposed to the first pole, and the first pole [30] receives the secondary cavity (second lumen [35]) (Figure 1B). It would have been obvious to a person having ordinary skill in the art at the time the invention was made to modify the cross-section of the catheter body, of the modified catheter of Frazee et al and Don Michael, with an oval cross-section, as taught by Prosl, as an obvious design choice to the user, since regardless of the cross-sectional shape of the catheter, the catheter will function to perfuse a substance into a vessel.

8. Claims 9-11 are rejected under 35 U.S.C. 103(a) as being unpatentable over Frazee et al and Don Michael, as applied to claim 1 above, and further in view of Petersen (US 3,490,457).

In regards to claim 9, in a modified catheter of Frazee et al and Don Michael, Frazee et al teaches that said second occluding means comprises an occluding body [127], suitable for being introduced into said main cavity [101]; however, Frazee et al does not teach that an insertion

cable is firmly connected to said occluding body [127]. Petersen teaches a catheter (Figure 1) having a second occluding means comprising an occluding body (obturator tip [50]) and an insertion cable (handle [51]). It would have been obvious to a person having ordinary skill in the art at the time the invention was made to modify the second occluding means, of the modified catheter of Frazee et al and Don Michael, with an insertion cable, as taught by Petersen, as the insertion cable will allow the user the ability to control the depth of the occluding body within the catheter.

In regards to claims 10 and 11, in a modified catheter of Frazee et al, Don Michael, and Petersen, neither Frazee et al nor Petersen teach that the occluding body is substantially spherical or frustoconical in shape, since Frazee et al teaches that the occluding body [127] is cylindrical (Figure 7), and Petersen teaches that the occluding body [50] is conical (Figure 2)(column 3, lines 13-16). However, at the time the invention was made, it would have been an obvious matter of design choice to a person having ordinary skill in the art to modify the shape of the occluding body to be either spherical or frustoconical because Applicant has not disclosed that a spherical or frustoconical occluding body provides an advantage, is used for a particular purpose, or solves a stated problem. One of ordinary skill in the art, furthermore, would have expected Applicant's invention to perform equally well with a cylindrical or conical occluding body because regardless of the shape of the occluding body, the occluding body will function to at least partially occlude the main cavity to minimize fluid flow towards the open distal end of the catheter.

9. Claim 19 is rejected under 35 U.S.C. 103(a) as being unpatentable over Frazee et al and Don Michael, as applied to claim 18 above, and further in view of Zhang (US 5,971,958).

In regards to claim 19, in a modified catheter of Frazee et al and Don Michael, Frazee et al does not teach that said main pathway [103] comprises a threaded section capable of producing a threaded connection with a corresponding threaded portion of said second occluding means [127]. Zhang teaches a catheter with a main pathway (introducer hub, *not referenced*) comprising a threaded section capable of producing a threaded connection with a corresponding threaded portion of a second occluding means (obturator, *not referenced*) (column 10, lines 23-65). It would have been obvious to a person having ordinary skill in the art at the time the invention was made to provide a threaded section on the main pathway and a threaded portion on the second occluding means, of the modified catheter of Frazee et al and Don Michael, so that a threaded engagement, of the main pathway and the second occluding body, will inhibit the rotational disengagement of the main pathway and the second occluding body (column 10, lines 23-65).

### ***Conclusion***

10. The prior art made of record and not relied upon is considered pertinent to applicant's disclosure: Ouriel et al (US 6,755,813), Parker et al (US 5,389,074) and Jackson (US 4,850,969).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to SHEFALI D. PATEL whose telephone number is (571) 270-3645. The examiner can normally be reached on Monday through Thursday from 8am-5pm Eastern time.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Kevin C. Sirmons can be reached on (571) 272-4965. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Shefali D Patel/  
Examiner, Art Unit 3767  
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